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(54) **SYSTEM FOR ENDOLUMINAL GRAFTING OF BIFURCATED OR BRANCHED VESSELS**

**SYSTEM ZUM ENDOLUMINALEN TRANSPLANTIEREN VON GEFÄSSEN MIT BIFURKATION
ODER ABZWEIGUNG**

**SYSTEME RELATIF A LA GREFFE ENDOLUMINALE DE VAISSEaux A DEUX BRANCHES OU
RAMIFIES**

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(73) Proprietor: **Medtronic AVE, Inc.**
Santa Rosa, California 95403 (US)

(72) Inventor: **Wisselink, Willem**
8251 AL Dronten (NL)

(74) Representative: **Whitaker, Iain Mark et al**
Sommerville & Rushton
Business Link Building
45 Grosvenor Road
St. Albans, Hertfordshire AL1 3AW (GB)

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EP-A- 0 792 627 **WO-A-95/21592**
WO-A-97/12562 **DE-A- 19 533 589**
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Description

[0001] The present invention relates generally to medical devices and methods, and more particularly to a system and method for endoluminal grafting of blood vessels or other tubular anatomical conduits which have furcations or side branches extending therefrom.

Background of the Invention

[0002] Endoluminal grafting is a relatively noninvasive method for placing a tubular graft within the lumen of an anatomical conduit, such as a blood vessel. In certain cardiovascular applications of the technique, an endovascular graft may be implanted within an aneurysmic segment of a blood vessel (i.e., a blood vessel which has a region of localized dilatation wherein the blood vessel wall has become distended and weakened) to form a prosthetic flow conduit through the aneurysm, and to effectively isolate weakened portion of the blood vessel wall from the hemodynamic forces and pressures of the flowing blood.

[0003] The prior art has included numerous endovascular grafts of varying design. In general, these endovascular grafts typically comprise a tube of pliable material (e.g., expanded polytetrafluoroethylene (ePTFE) or woven polyester) in combination with a graft anchoring component (e.g., a stent, a frame, a series of wire rings, hooks, barbs, clips, staples, etc.) which operates to hold the tubular graft in its intended position within the blood vessel. Most commonly, the graft anchoring component is formed of a radially expandable stent or frame which is either a) incorporated into the body of the tubular graft or b) formed separately from the graft and deployed within the graft lumen, and which is expandable to exert outwardly directed radial pressure against the surrounding blood vessel wall—thereby frictionally holding the graft in place. In operation, endovascular grafts which incorporate radially expandable graft anchoring devices are initially disposed in a radially collapsed configuration which is sufficiently compact to allow the graft to be transluminally advanced through the vasculature until it reaches the intended site of implantation. Thereafter, the graft (and the accompanying graft anchoring device) expands to a radially expanded configuration which is large enough to exert the desired outwardly-directed pressure against the blood vessel wall. In some embodiments, hooks, barbs, or other projections formed on the graft anchoring device, will insert into the wall of the blood vessel to ensure that the graft will be firmly held in its desired position, without slipping or migrating after implantation. These radially expandable graft anchoring devices are generally classifiable as either a.) self-expanding or b) pressure-expandable. Graft anchoring devices of the "self-expanding" are usually formed of a resilient material (e.g., spring metal) or shape memory alloy which automatically expands from a radially collapsed configuration to a radially expanded

configuration, when relieved of surrounding constraint (e.g., a surrounding tubular sheath or catheter wall). On the other hand, those of the "pressure-expandable" variety are typically formed of malleable wire or other plastically deformable material which will deform to a radially expanded configuration in response to the exertion of outwardly directed pressure thereagainst—as by inflation of a balloon or actuation of another pressure-exerting apparatus which has been positioned within the graft anchoring device.

[0004] Endovascular grafting is a clinically viable alternative to traditional surgery in patients who suffer from aneurysms of the aorta. Indeed, many patients who are diagnosed with aortic aneurysms are in relatively poor health, and may be characterized as poor surgical risks. Moreover, the traditional surgical approach to repair of aortic aneurysms requires cross-clamping of the aorta above the aneurysm, which can result in ischemic damage to organs or other untoward results. Nonetheless, if allowed to remain untreated, a substantial percentage of aortic aneurysms may ultimately dissect or rupture, with catastrophic consequences. Thus, endovascular grafting offers a potential means for repair of aortic aneurysms, without the risks and potential complications of traditional aneurysm repair surgery.

[0005] Depending on which region(s) of the aorta is/are involved, the aneurysm may extend into areas of bifurcation (i.e., the inferior end of the aorta where it bifurcates into the iliac arteries) or segments of the aorta from which smaller "branch" arteries extend. In this regard, the various types of aortic aneurysms may be classified on the basis of the region(s) of aneurysmic involvement, as follows:

A. Thoracic Aortic Aneurysms:

- Aneurysms involving the ascending thoracic aorta
- Aneurysms involving the aortic arch and branch arteries which emanate therefrom (i.e., the subclavian arteries)

B. Thoracoabdominal Aortic Aneurysms:

- Aneurysms involving the descending thoracic aorta and branch arteries which emanate therefrom (i.e., thoracic intercostal arteries) and/or the suprarenal abdominal aorta and branch arteries which emanate therefrom (i.e., renal, superior mesenteric, celiac and/or intercostal arteries).

C. Abdominal Aortic Aneurysms:

- Aneurysms involving the pararenal aorta and the branch arteries which emanate therefrom (i.e., the renal arteries)

■ Aneurysms involving the infrarenal aorta with or without iliac involvement

[0006] Unfortunately, not all patients diagnosed with aortic aneurysm are presently considered to be candidates for endovascular grafting. This is largely due to the fact that most of the endovascular grafting systems of the prior art are not designed for use in regions of the aorta from which side branches (i.e., carotid, innominate, subclavian, intercostal, superior mesenteric, celiac or renal arteries) extend. In fact, most of the clinical experience with endoaortic grafting has been for the treatment of infrarenal aneurysms, with or without iliac involvement. Examples of endovascular grafting methods and systems useable to treat such infrarenal aneurysms include those described in the following United States patents: 4, 577, 631 (Kreamer); 5, 211, 658 (Clouse); 5, 219, 355 (Parodi et al.); 5, 316, 023 (Palmaz et al.); 5, 360, 443 (Barone et al.); 5, 425, 765 (Tifenbrun et al.); 5, 609, 625; (Piplani et al.); 5, 591, 229 (Parodi et al.); 5, 578, 071 (Parodi); 5, 571, 173 (Parodi); 5, 562, 728 (Lazarus et al.); 5, 562, 726 (Chuter); 5, 562, 724 (Vorwerk et al.); 5, 522, 880 (Barone et al.); and 5, 507, 769 (Marin et al.).

[0007] Most if not all of the endovascular grafts which have been designed for use in treating infrarenal aneurysms require that a proximal "neck" (e.g., at least two (2) centimeters of non-aneurysmic aorta) exist inferior to the renal arteries, in order to provide a region where the superior end of the graft may be securely anchored in place, without blocking or restricting blood flow into the renal arteries. The deployment of endovascular grafts within regions of the aorta from which branch arteries extend (e.g., regions of the aorta from which the renal, superior mesenteric, celiac, intercostal, and/or subclavian arteries emanate) present additional technical challenges because, in those cases, the endovascular graft must be designed, implanted and maintained in a manner which does not impair the flow of blood into the branch arteries.

[0008] United States Patent No. 5, 425, 765 (Tifenbrun et al.) has described an endovascular graft which has one or more openings or fenestrations formed at specific locations, to allow blood to flow from the aorta into one or more branch arteries. However, such fenestrations do not form discrete conduit(s) through which blood is channeled into each branch artery. As a result, the area surrounding the fenestrations could be prone to i) the leakage of blood into the space between the outer surface of the aortic graft and the surrounding aortic wall or ii) post-implantation migration or movement of the graft causing misalignment of the fenestration(s) and the branch artery(ies)—with resultant impairment of flow into the branch artery(ies).

[0009] Thus, in view of the above-discussed limitations and shortcomings, there remains a need in the art for the development of new endovascular grafting systems and methods which a) may be useable for en-

dovascular grafting in regions of a blood vessel (e.g., aorta) from which branch blood vessels (e.g., carotid, innominate, subclavian, intercostal, superior mesenteric, celiac, renal or iliac arteries) extend, and/or b) may enable more aortic aneurysm patients to be considered as candidates for endovascular repair, and/or c) may otherwise advance the state of the art of endovascular grafting to improve patient outcomes or lessen complications.

[0010] A bifurcated endoluminal graft for deployment in an anatomical conduit, having a branch graft opening as well as a first connector member and a tubular branch graft with a second connector member is disclosed in PCT/WO 95/21592. the claims hereinafter are characterized over this prior art.

Summary of the Invention

[0011] The present invention provides a system for endoluminal grafting of a blood vessel or other anatomical conduit, in a region where one or more branch anatomical conduit(s) (e.g., side branches, furcations, etc.) extend from the anatomical conduit.

[0012] In accordance with the present invention, there is provided an endoluminal grafting system which comprises:

(a) a tubular primary graft for deployment in the anatomical conduit;

(i) said primary graft having first and second ends and a lumen extending longitudinally therethrough.

(ii) a branch graft opening formed at a first location in the primary graft;

(iii) a first connector member formed adjacent the branch graft opening; and,

(iv) one or more primary graft anchoring components operative to hold at least the first and second ends of the primary graft in substantially abutting contact with the anatomical conduit; and

(b) a tubular branch graft for deployment into the branch anatomical conduit;

(i) said branch graft having a proximal end, a distal end, and a lumen extending longitudinally therethrough;

(ii) a second connector member being associated with the proximal end of the branch graft, said second connector member being configured to engage said first connector as the proximal end of the branch graft is brought into approximation with the branch graft opening of the primary graft, wherein the one or more primary graft anchoring components hold the primary graft at locations on either side of said branch

anatomical conduit, said primary graft anchoring components being thereby operative to hold the primary graft in a substantially fixed position within the lumen of the anatomical conduit such that its branch graft opening is substantially aligned with said branch anatomical conduit; and said branch graft being advanceable distal-end-first through the branch graft opening and into the branch anatomical conduit; said second connector member being configured to automatically frictionally engage said first connector to form a connection between the proximal end of the branch graft and the primary graft.

[0013] In some applications, it will be desirable for the branch graft component of the above-summarized endoluminal grafting system to further include, or to be used in conjunction with;

iii) a branch graft anchoring device (e.g., a radially expandable stent, frame, rings, and/or an adhesive, suture(s), staple(s), etc.) which is operative to hold at least the distal end of the branch graft in contact with the surrounding wall of the branch anatomical conduit.

[0014] The above-summarized endoluminal grafting system may be implanted within the branched anatomical conduit by a method which comprises the following procedural steps:

a) transluminally advancing the primary graft into the branched anatomical conduit;
 b) positioning the primary graft within the anatomical conduit such that the branch opening is aligned with the branch anatomical conduit;
 c) utilizing the primary graft anchoring device to anchor the primary graft within the anatomical conduit;
 d) transluminally advancing the branch graft into the lumen of the primary graft;
 e) passing the distal end of the branch graft through the branch opening and into the branch anatomical conduit; and,
 f) advancing the branch graft into the branch anatomical conduit until the second connector on the proximal end of the branch graft engages the first connector of the primary graft, thereby connecting the branch graft to the primary graft such that body fluid (e.g., blood) may flow from the lumen of the primary graft, through the branch opening, and through the lumen of the branch graft.

[0015] The above summarized method may optionally include step of:

g) utilizing a branch graft anchoring device to anchor at least the distal end of the branch graft to the surrounding wall of the branch anatomical conduit.

[0016] A method for implanting a modular, bifurcated endoluminal graft within a bifurcated anatomical conduit (i.e., a conduit having a main portion (e.g., aorta), a first furcation (e.g., right iliac) and a second furcation (e.g., left iliac), through a single access opening (e.g., cut down incision or percutaneous puncture site) suitably generally comprises the steps of:

a. providing a modular endoluminal graft which comprises i) a primary graft which is configured to be positioned in the main portion and first furcation of the anatomical conduit, and which has an opening formed therein, and ii) a branch graft which has a proximal end and a distal end and which is configured to be positioned in the second furcation of the anatomical conduit with its proximal end connected to the opening formed in the main graft;
 b. forming an access opening which leads into the first furcation of the anatomical conduit;
 c. advancing the bifurcated primary graft through the access opening and positioning the primary graft in the main portion and first furcation of the anatomical conduit;
 d. advancing a curved guide catheter having an internal angle of curvature less than 90 degrees, into the main portion of the primary graft;
 e. advancing a guidewire having a distal end, through the curved guide catheter, such that the distal end of the guidewire passes out of the curved guide catheter, through the opening formed in the primary graft, and into the second furcation of the anatomical conduit;
 f. advancing a catheter having the branch graft mounted thereon, over the guidewire until the distal end of the branch graft extends into the second furcation of the anatomical conduit and the proximal end of the branch graft is connected to the opening formed in the primary graft; and,
 g. removing said guidewire and said catheter.

In some applications of this method, an anchorable guidewire may be inserted in step e and the guidewire will then be anchored in the second furcation of the anatomical conduit, to prevent the guidewire from being inadvertently pulled out of the second furcation during the procedure. In procedure wherein such anchorable guidewire is used, the anchoring apparatus of the guidewire will be actuated prior to step f and will be deactivated prior to step g.

[0017] By way of example, there is provided a new anchorable guidewire system which is useable in conjunction with the above-summarized method for implanting a bifurcated endoluminal graft and/or with various other medical or surgical procedures. This anchorable guidewire system comprises:

a) a balloon-anchorable guidewire which compris-

es:

- i) an elongate guidewire having a proximal end and a distal end;
- ii) a balloon formed on the guidewire at a first location;
- iii) a lumen which extends longitudinally through the guidewire from its proximal end to said balloon, said lumen being useable to pass inflation fluid into and out of said balloon; and,

b) an inflation syringe/lumen-plugging apparatus which comprises:

- i) a syringe having a plunger and a barrel, said syringe being connectable to the proximal end of the guidewire such that when the plunger of said syringe is advanced in a first direction it will force inflation fluid through said lumen in the distal direction and into said balloon and, when said plunger is retracted in a second direction it will draw inflation fluid out of said balloon and through said lumen in the proximal direction; and,
- ii) a lumen-plugging member releasably mounted on the plunger of the syringe such that, when the plunger is advanced in the first direction to a predetermined point, the lumen-plugging member will become inserted into the guidewire lumen and is subsequently releasable from the plunger, said lumen-plugging member being thereby operative to prevent inflation fluid from escaping from said lumen when the syringe is removed.

[0018] In operation, the balloon-anchorable guidewire is inserted to its desired position and the inflation syringe/lumen-plugging apparatus is attached to its proximal end. The syringe plunger is then advanced in the distal direction, thereby inflating the balloon and anchoring the distal end of the guidewire at a desired position within the body. After the plunger has been advanced far enough to cause the lumen blocking member to be inserted into the proximal end of the guidewire lumen, the lumen blocking member is released (e.g., broken away or disconnected) from the plunger, and the lumen blocking member is allowed to remain inserted in proximal end of the lumen-acting as a plug to prevent fluid from escaping from the lumen. The syringe may then be detached and removed from the proximal end of the guidewire, and one or more catheters or other transluminal devices may be advanced and/or retracted over the proximal end of the guidewire. Thereafter, when it is desired to remove the guidewire, the exteriorized portion of the guidewire is cut (e.g., transected by a pair of standard scissors), thereby allowing the inflation fluid to leak from the balloon and out of the guidewire lumen. After the balloon has been deflated in this manner, the

guidewire may be extracted and removed from the body.

[0019] Further aspects and details of the present invention will become apparent to those skilled in the art upon reading and understanding of the following detailed description and the drawings to which it refers.

Brief Description of the Drawings

[0020]

Figure 1 is a schematic showing of a human body having an endoluminal grafting system of the present invention implanted to repair an abdominal aortic aneurism which involves the renal arteries. Figure 1a is an enlarged view of the perirenal aorta and endovascular grafting system shown in Figure 1.

Figure 1b is a partial longitudinal sectional view of an endoluminal grafting system of the present invention, showing a preferred means for connecting the proximal end of the branch graft to the primary graft.

Figure 1c is an exploded view of the graft components shown in Figure 1b.

Figure 1d shows a first step in a preferred method for placing the branch graft(s) of an endoluminal grafting system of the present invention after the primary graft has been implanted.

Figure 1e shows a second step in a preferred method for placing the branch graft(s) of an endoluminal grafting system of the present invention after the primary graft has been implanted.

Figure 1f is a schematic showing of a portion of a human aorta having an endoluminal grafting system of the present invention implanted to repair a thoracoabdominal aortic aneurysm which involves the renal, superior mesenteric and celiac arteries. Figure 2 is a schematic showing of a human body having a the primary graft component of an endoluminal grafting system of the present invention implanted in the distal aorta and one iliac artery, to begin the repair of an infrarenal abdominal aortic aneurism which has bilateral iliac involvement

Figure 2a is a view of the aneurysmic portion of the aorta in Figure 2, showing a first step in a preferred method for placing an iliac branch graft in the remaining iliac artery to complete the implantation of the endoluminal grafting system of the present invention.

Figure 2b is a view of the aneurysmic portion of the aorta in Figure 2a, showing a second step in a preferred method for placing an iliac branch graft in the remaining iliac artery to complete the implantation of the endoluminal grafting system of the present invention.

Figure 2c is a view of the aneurysmic portion of the aorta in Figure 2b, following placement of an iliac branch graft in the remaining iliac artery and

complete implantation of the endoluminal grafting system of the present invention.

Figure 3 is a schematic showing of an infrarenal aortic aneurysm having bilateral iliac involvement and wherein the primary graft component of an endoluminal grafting system of the present invention has been implanted within the distal aorta and one iliac artery, to begin the repair of the aneurism, and a preferred balloon-anchorable guidewire of the present invention is being advanced into the contralateral iliac artery to facilitate placement of a branch graft within the contralateral iliac artery.

Figure 3a shows the proximal end of the preferred anchorable guidewire of Figure 3 connected to an inflation syringe/lumen plugging apparatus, prior to inflation of the guidewire balloon.

Figure 3b shows the proximal end of the preferred anchorable guidewire of Figure 3 connected to an inflation syringe/lumen plugging apparatus, following inflation of the guidewire balloon but before detachment of the inflation syringe.

Figure 3c shows the proximal end of the balloon anchorable guidewire of Figure 3 after inflation of the guidewire balloon and detachment of the inflation syringe, with the lumen-plugging apparatus having been separated from the inflation syringe and remaining in place within the proximal end of the guidewire lumen.

Detailed Description of the Preferred Embodiments

[0021] Referring now to the drawings, wherein the showings are for the purpose of illustrating presently preferred embodiments of the invention only and are not intended to limit the scope of the invention in any way.

[0022] The accompanying figures include various showings of human anatomical structures, and such anatomical structures are labeled according to the following legend:

Aorta	A
Celiac Artery	CA
Femoral Artery	F
Heart	H
Iliac Arteries	IL (IL ₁ and IL ₂)
Kidneys	K
Renal Arteries	RA
Superior Mesenteric Artery	SMA

A. General Structure and Function of Endoluminal Grafting Systems For Branched Anatomical Conduits

[0023] As shown generally in Figures 1-2c, each preferred endoluminal grafting system 10 of the present invention generally comprises a primary graft 12 having at least one branch graft opening 14 formed therein, and

at least one branch graft 16 which is advancable out of the branch graft opening 14 and into a branch anatomical conduit (e.g., a side branch or furcation of a blood vessel). A branch graft connector apparatus 17 is incorporated into the primary graft 12 and/or branch graft(s) 16 to connect the proximal end of the branch graft(s) 16 to the primary graft 12. One or more primary graft anchoring devices 18, such as radially expandable stent(s) or frame(s), is/are used to frictionally hold the primary graft 12 in a fixed position in the lumen of the primary anatomical conduit (e.g., aorta). Also, where necessary, one or more branch graft anchoring devices 20, such as radially expandable stent(s) or frame(s), may be used to frictionally hold at least the distal end of the branch graft(s) 16 in fixed position in the lumen(s) of the branch anatomical conduit(s) (e.g., carotid, innominate, subclavian, intercostal, superior mesenteric, celiac, renal or iliac arteries). These primary graft anchoring device(s) 18 and branch graft anchoring device(s) 20 may be formed or incorporated into the body of the primary and branch grafts 12, 16 or, alternatively, they may be formed as separate structures (e.g., separate self-expanding or pressure-expandable stents) which are positioned within the lumens of the primary and branch grafts 12, 16, to accomplish the desired anchoring of the primary and branch grafts 12, 16, as shown.

[0024] Each branch graft opening 14 of the primary graft 12 is strategically placed, and preferably reinforced and marked for radiographic visualization, to facilitate precise alignment of each branch graft opening 14 with the particular branch anatomical conduit (e.g., carotid, innominate, subclavian, intercostal, superior mesenteric, celiac, renal or iliac arteries) into which its branch graft 16 is to extend. The desired strategic placement of each branch graft opening 14 may be facilitated by custom-forming the branch graft opening(s) 14 in the primary graft 12, prior to implantation of the primary graft 12. Such pre-implantation, custom formation of the branch graft opening(s) 14 may be accomplished through the use of spiral computed tomography data of the vascular anatomy of the particular patient in whom the endovascular grafting system 10 is to be implanted. Notably, in applications where the primary graft 12 is being prepared for implantation in the thoracic or thoraco-abdominal aorta, the branch graft openings 14 may be sized and positioned to align with relatively small side branch vessels (e.g., carotid, innominate, subclavian, intercostal, superior mesenteric, celiac, or renal arteries) which, unlike the iliac bifurcation at the inferior end of the aorta, require a branch graft 16 be passable through such branch graft opening(s) 14 and into the smaller side branch vessel (e.g., carotid, innominate, subclavian, intercostal, superior mesenteric, celiac, or renal arteries) at an angle of approximately 80-90 degrees relative to the axis of the aorta.

[0025] After the branch graft opening(s) 14 have been formed at the desired locations (and the appropriate portions of the branch graft connector 17 have been installed

into the primary graft adjacent the branch graft opening (s) 14), the primary graft 12 is inserted via an introducer into the femoral artery, and advanced under radiographic guidance into the site (e.g., the aneurysmic segment of vessel) where the primary graft 12 is to be implanted. This insertion of the primary graft 12 (and any primary graft anchoring device(s) 18 which are formed separately from the graft 12) may be facilitated by mounting the primary graft 12 (and any such separate graft anchoring devices 18) on a balloon catheter or other suitable delivery catheter capable of carrying the primary graft (and any separate graft anchoring device(s) 18) to the intended site of implantation. The primary graft 12 and any separate primary graft anchoring device 18 is/are then radially expanded or otherwise deployed such that the primary graft becomes anchored in a substantially fixed position within the primary anatomical conduit (e.g., aorta A). In applications wherein the endoluminal grafting system 10 is being used to repair an aneurysm, the hemodynamic pressure within the aneurysm sac (i.e., the space within the aneurysm but outside the primary graft 12), as well as the hemodynamic pressure within the branch anatomical conduit(s) which emanate from the aneurysmic segment of vessel, will be substantially unaffected by the initial placement of the primary graft 12 because, until such time as the branch graft(s) 14 have been introduced, blood will continue to flow out of the branch graft opening(s) 14 of the primary graft 12.

[0026] After the primary graft 12 has been positioned and anchored within the primary anatomical conduit (e.g., aorta A), a guidewire 30 may be transluminally advanced through the lumen of the primary graft 12, out of a branch opening 14 and into the branch anatomical conduit (e.g., In some applications, a guide catheter may be introduced into the lumen of the primary graft 12 to facilitate passage of the guidewire 30 out of the desired branch graft opening 14.

[0027] A branch graft 16 is then mounted on a balloon catheter 32, and the distal end of the branch graft may be drawn taught about the catheter balloon 34 by a purse string suture 36 (e.g., 7.0 polypropylene suture material). The balloon catheter 32 having the branch graft 16 mounted thereon is then advanced over the guidewire 30 until the proximal end of the branch graft 16 becomes connected to the branch graft opening 14 of the primary graft 12 by way of the branch graft connector apparatus 18. Thereafter, the catheter balloon 34 is inflated, causing the purse string suture 36 to break and the distal end of the branch graft 16 to radially expand into contact with the surrounding wall of the branch anatomical conduit.

[0028] A presently preferred branch graft connector apparatus 17 is shown in detail in figures 1b-1e. The components of this branch graft connector apparatus 17 associated with the primary graft 12 comprise: a first rigid or semirigid ring 40 which surrounds the branch graft opening 14, and a rigid or semirigid frustoconical member 42 which extends outwardly from the first ring mem-

ber 40, as shown in Figure 1b. The components of this branch graft connector apparatus associated with the branch graft 16 comprise a rigid or semirigid, tapered proximal portion 44, a second rigid or semirigid ring member 46 on the proximal end of the tapered proximal portion 44, and a third rigid or semirigid ring member 48 formed about the tapered proximal portion 44 at a spaced distance from the second ring member. The distance between the outer surface OS of the second ring member 46 and the inner surface IS of the third ring member 48 is substantially the same as the distance between the inner surface IS of the first ring member 40 and the distal end DE of the frustoconical member 42. In this manner, as the branch graft 12 is advanced, distal end first, out of the branch graft opening 14 it will reach a point of maximum advancement whereat the inner surface IS of the first ring member 40 will abut against the outer surface OS of the second ring member 46, and the distal end DE of the frustoconical member 42 will abut against the inner surface IS of the third ring member 48. This will create a substantially fluid-tight seal between the proximal end of the branch graft 16 and the body of the primary graft 12. Additionally, this may form a snap-fit connection which will prevent the branch graft 14 from slipping or undergoing inadvertent retraction back into the lumen of the primary graft 12.

[0029] It will be appreciated that an audible or tactilely discernible indicator (e.g., a "snap") may occur as the separate components of the branch graft connector 17 come together, thereby indicating to the operator that the desired engagement and sealing of the proximal end of the branch graft 16 to the primary graft 12 has been accomplished. Also, the ring members 40, 46, and 48 may be formed of radiographically visible material to enable the operator to radiographically confirm that the branch graft 16 has been firmly connected to and engaged with the primary graft 12. Additionally, it is to be appreciated that the branch graft connector 17 need not necessarily be a mechanical connector apparatus as shown in the drawings, but may alternatively comprise any mechanical, electromagnetic, frictional, or chemical (e.g., adhesive, coupling, or fusing agent) capable of joining the proximal end of the branch graft 16 to the primary graft 12 in substantially fluid-tight sealing contact.

B. A Specific Endoluminal Grafting System and Method for Repair of Aortic Aneurysms Having Iliac Involvement

[0030] Figures 2-3c are specifically directed to embodiments of the endovascular grafting system 10a for the repair of aneurysms of the infrarenal aorta A which extend into the iliac arteries IL. This embodiment of the endoluminal grafting system 10a differs from the above-described embodiment 10 in the primary graft 21a has a bifurcation 64 formed therein, and includes a main aortic portion 60 and one iliac portion 62. The branch graft

opening 14 of this embodiment 10a is formed on one side of the bifurcation 64 to facilitate passage of the branch graft 14 into the iliac artery IA which is opposite the iliac artery IA into which the iliac portion 62 of the primary graft 12a has been positioned. Also, in the primary graft component 12a of this embodiment 10a the primary graft anchoring devices 18 are located at the superior end of the main portion 60 and the inferior end of the iliac portion 62, as shown in Figure 2a.

[0031] With reference to the step-wise showings of Figures 2a-2c, a method for implanting the bifurcated endoluminal graft of the present invention comprises the steps of:

1. Forming an access opening 66 into one of the femoral arteries F, and advancing a guidewire 30 through such access opening 66, through the contiguous iliac artery IL₁, and into the aorta A;
2. Advancing the bifurcated primary graft 12a over the guidewire 30, and positioning the primary graft 12a in the aorta A and first iliac artery IL₁, such that its branch graft opening 14a is directed toward the other iliac artery IA₂, as shown in Figure 2a;
3. Causing the primary graft anchoring device 18 to hold at least the superior end of the aortic portion 60 of the primary graft 12a in the aorta A, and the inferior end of the iliac portion 62 of the primary graft 12a in the first iliac artery IA₁;
4. Advancing a curved guide catheter 68, having an angle of curvature AC less than 90 degrees, into the lumen of the aortic portion 60 of the primary graft 21 a;
- 5 Advancing the guidewire 30 through the guide catheter 68, such that the distal end of the guidewire 30 passes out of the distal end of the curved guide catheter 68 and into the other iliac artery IA₂;
6. Advancing the balloon catheter 32 having the branch graft 14 mounted thereon, over the guidewire 30 as shown in Figure 2b, until the branch graft connector apparatus 17 becomes connected, thereby forming a fluid tight sealing connection between the proximal end of the branch graft 14 and the primary graft 12a;
7. Inflating the balloon 34 of the balloon catheter 32 to break the purse string stitch 36 and to radially expand the branch graft anchoring device 20 (if present);
8. Deflating the catheter balloon 32; and,
9. Removing the guidewire 30 and balloon catheter 32 from the body.

[0032] Thus, by the above described steps of the method of the present invention, an endoluminal grafting system 10a may be implanted in a bifurcated anatomical conduit, with only one (1) access opening (e.g., puncture site or incision) formed in the body.

[0033] In at least some applications of the method it may be desirable to anchor the distal end of the

guidewire 30 within the contralateral iliac artery IL₂ to prevent the guidewire from becoming displaced (i.e., pulled out of the contralateral iliac artery IL₂ during the advancement of the branch graft 14. Various types of anchorable (e.g., balloon-anchorable or tip-deflectable) guidewires could be useable in (or modifiable for use in) this procedure. Examples of such previously known anchorable guidewires include those described in United States Patent Nos. 5, 167, 239 (Cohen et al.); 5, 480, 362 (Hammerslag et al.) and 5, 497, 782 Fugoso).

[0034] By way of example an anchorable guidewire system 78 as shown in Figures 3-3c is described more fully herebelow

15 C. A PREFERRED ANCHORABLE GUIDEWIRE SYSTEM

[0035] Figures 3-3c show an anchorable guidewire system 78. This anchorable guidewire system 78 is useable in the performance of the above-described method for repairing aneurysms of the aorta which have iliac involvement. Additionally, this anchorable guidewire system 78 may have numerous other applications in the performance of various medical procedures in interventional cardiology, interventional radiology, gastroenterology, and surgery.

[0036] The preferred anchorable guidewire system 78 comprises a pliable guidewire 80 and an inflation syringe/lumen plugging apparatus 82. The guidewire 80 has a balloon 83, such as a soft shape-conformable balloon (e.g. latex), formed about the distal portion of the guidewire 80 and a balloon inflation/deflation lumen 81 extending longitudinally through the guidewire 80 to permit balloon inflation fluid (e.g., saline, CO₂, room air) to be passed into and out of the balloon. The inflation syringe/lumen plugging apparatus 82 comprises, a syringe having a generally cylindrical barrel 85 and a plunger 86 which is in fluid-tight contact with the barrel 85. The plunger 86 slides longitudinally back and forth within the barrel 85 to alternately a) draw fluid into the barrel 85, and b) expel fluid out of the barrel 85. A lumen-plugging apparatus 84 is attached to the distal face of the plunger 86 and protrudes distally therefrom, as shown. This lumen-plugging apparatus is detachable from the plunger 86 by any acceptable means, such as a breakable attachment which will be severed when the plunger 86 is rotated while the lumen-plugging apparatus 84 is held in a fixed non-rotating position. To facilitate this, the lumen plugging apparatus 84 may, when inserted into the proximal end of the guidewire lumen 81, engage the surrounding wall of the guidewire lumen 81 tightly enough to prevent the lumen plugging apparatus 84 from rotating when the plunger 86 is rotated. In this rotation-preventing surface texturing, such as a system of longitudinal ribs or corresponding longitudinal ribs & grooves (not shown), may be formed on the outer surface of the lumen plugging apparatus 84 and/or the surrounding wall of the guidewire lumen 81 such that rota-

tion of the lumen plugging apparatus 84 will be frictionally prevented or deterred.

[0037] As shown in Figure 3a the inflation syringe/lumen plugging apparatus 82 is initially attached to the guidewire 80, with its plunger 86 in a withdrawn position, and with the proximal end of the guidewire inserted into the barrel 85 of the syringe 82. In this manner,

[0038] inflation fluid is contained within the barrel 85 for use in inflating the balloon 83, when desired.

[0039] Thereafter, as shown in figure 3b, when it is desired to inflate the balloon 83, the plunger 86 of the syringe 82 is advanced in the distal direction, thereby forcing inflation fluid out of the barrel 85, through the guidewire lumen 81, and into the balloon 83. This results in inflation of the balloon 83 and anchoring of the distal end of the guidewire 80. This also causes the lumen plugging apparatus 84 on the distal face of the plunger 86 to be forced into the proximal end of the guidewire lumen 81, thereby plugging the lumen 81 to prevent the balloon inflation fluid from escaping from the balloon. Also, in the particular embodiment shown in the drawings, this will also cause the lumen plugging apparatus 84 to be non-rotatably engaged to the guidewire 80.

[0040] Thereafter, when it is desired to pass the balloon catheter 32 or other device over the proximal end of the guidewire 80, the plunger 86 of the syringe 82 is rotated. Such rotation of the plunger 86 causes the lumen-plugging apparatus 84 to break away from, or separate from, the plunger 86. This allows the syringe 82 to be removed, as shown in Figure 3c, while the lumen plugging apparatus remains inserted within the proximal end of the guidewire lumen 81 to prevent the inflation fluid from escaping from the guidewire balloon 83.

[0041] Subsequently, when it is desired to deflate the balloon 83 to allow the guidewire to be extracted, the operator may simply cut the guidewire 80 at a location distal to the lumen plugging apparatus 84, thereby allowing the balloon inflation fluid to escape from the lumen 81 and causing the balloon 83 to deflate.

[0042] It will be appreciated that, although the invention has been described hereabove with reference to certain presently preferred embodiments or examples shown in the drawings, various modifications could be made to those presently preferred embodiments or examples without departing from the scope of the invention as claimed. Accordingly, it is intended that all such modifications to the above-described preferred embodiments or examples be included within the scope of the following claims.

Claims

1. A system for endoluminal grafting of an anatomical conduit (A) having at least one branch anatomical conduit (IA, RA) emanating therefrom, said system comprising:

(a) a tubular primary graft (12) for deployment in the anatomical conduit;

(i) said primary graft (12) having first and second ends and a lumen extending longitudinally therethrough.

(ii) a branch graft opening (14) formed at a first location in the primary graft (12);

(iii) a first connector member (40, 42) formed adjacent the branch graft opening (14); and,

(iv) one or more primary graft anchoring components (18) operative to hold at least the first and second ends of the primary graft (12) in substantially abutting contact with the anatomical conduit; and

(b) a tubular branch graft (16) for deployment into the branch anatomical conduit;

(i) said branch graft (16) having a proximal end, a distal end, and a lumen extending longitudinally therethrough;

(ii) a second connector member (44, 46, 48) being associated with the proximal end of the branch graft (16), said second connector member (44, 46, 48) being configured to engage said first connector as the proximal end of the branch graft is brought into approximation with the branch graft opening of the primary graft, **characterised in that** the one or more primary graft anchoring components (18) hold the primary graft (12) at locations on either side of said branch anatomical conduit, said primary graft anchoring components (18) being thereby operative to hold the primary graft (12) in a substantially fixed position within the lumen of the anatomical conduit such that its branch graft opening (14) is substantially aligned with said branch anatomical conduit; and said branch graft (16) being advanceable distal-end-first through the branch graft opening (14) and into the branch anatomical conduit; said second connector member (44, 46, 48) being configured to automatically frictionally engage said first connector to form a connection between the proximal end of the branch graft and the primary graft.

2. The system of claim 1, wherein at least one of said primary graft (12) and said branch graft (16) is formed at least partially of woven polyester.

3. The system of claim 1, wherein at least one of said primary graft (12) and said branch graft (16) is formed at least partially of expanded polytetrafluor-

oethylene.

4. The system of claim 1, 2 or 3 wherein the branch graft opening (14) is formed in the side of the primary graft (12).

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5. The system of claim 1, 2 or 3 wherein a bifurcation is formed at one end of the primary graft (12) and said branch graft opening (14) is formed in one side of that bifurcation.

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6. The system of claim 5 wherein the primary graft (12) further comprises a pliable tubular graft extension (62) which extends from the side of the bifurcation which is opposite the side in which the branch graft opening (14) is formed.

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7. The system of any preceding claim, wherein the first connector member (40, 42) comprises an annular groove formed at least part way around the branch graft opening (14) in the primary graft (12).

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8. The system of any preceding claim wherein the one or more primary graft anchoring components are selected from the group of graft anchoring apparatus and substances consisting of:

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at least one radially expandable stent which is separate from the primary graft (12);

at least one radially expandable stent which is integrated into the primary graft (12);

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at least one radially expandable frame which is separate from the primary graft;

at least one radially expandable frame which is incorporated into the primary graft;

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at least one projection which protrudes at least partially through the wall of the anatomical conduit;

an adhesive;

at least one suture;

and

at least one staple.

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9. The system of any preceding claim wherein the branch graft (16) portion of said system further comprises a branch graft anchoring component (20) for holding at least the distal end of the branch graft (16) in contact with the wall of the branch anatomical conduit.

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10. The system of claim 9, wherein the branch graft anchoring component is selected from the group of graft anchoring apparatus and substances consisting of:

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at least one radially expandable stent which is separate from a first pliable tube of the branch graft (16);

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at least one radially expandable stent which is integrated into a first pliable tube of the branch graft (16);

at least one radially expandable frame which is separate from a first pliable tube of the branch graft (16);

at least one radially expandable frame which is incorporated into a first pliable tube of the branch graft (16);

at least one projection which protrudes at least partially through the wall of the anatomical conduit;

an adhesive;

at least one suture;

and at least one staple.

11. The system of any preceding claim wherein the second connector member (44, 46, 48) on the branch graft (16) is configured to frictionally engage the first connector member (40, 42) of the primary graft 12, as the branch graft (16) is advanced out the branch graft opening (14) of the primary graft (12).

12. The system of claim 11, wherein the first connector member (40, 42) on the primary graft (12) is an annular groove, and the second connector member (44, 46, 48) on the branch graft (16) is an annular ridge which is sized and configured to seat within and frictionally engage said annular groove.

13. The system of any preceding claim wherein at least a portion of the branch graft (16) is tapered so that its distal end is smaller in diameter than its proximal end.

14. The system of claim 13 wherein a portion of the branch graft (16), adjacent its proximal end, is frustoconical in configuration.

15. The system of claim 14, wherein the first connector member (44, 46, 48) is formed on the proximal end of the frustoconical segment (44) of the branch graft (16).

Patentansprüche

1. System zum endoluminalen Transplantieren einer anatomischen Röhre (A) mit wenigstens einer von derselben ausgehenden anatomischen Zweigröhre (IA, RA), wobei das System folgendes umfaßt:

(a) ein röhrenförmiges Haupttransplantat (12) zum Entfalten in der anatomischen Röhre,

(i) wobei das Haupttransplantat (12) ein erstes und ein zweites Ende und einen in Längsrichtung durch denselben verlaufen-

den Hohlraum,

(ii) eine an einer ersten Stelle im Haupttransplantat (12) geformte Zweigtransplantatöffnung (14),

(iii) ein erstes, angrenzend an die Zweigtransplantatöffnung (14) geformtes, Anschlußelement (40, 42) und

(iv) eine oder mehrere Haupttransplantat-Verankerungskomponenten (18), wirksam zum Halten wenigstens des ersten und des zweiten Endes des Haupttransplantats (12) in einem wesentlich anstoßenden Kontakt mit der anatomischen Röhre, hat, und

(b) ein röhrenförmiges Zweigtransplantat (16) zum Entfalten in die anatomische Zweigröhre,

(i) wobei das Zweigtransplantat (16) ein proximales Ende, ein distales Ende und einen in Längsrichtung durch denselben verlaufenden Hohlraum hat,

(ii) ein zweites Anschlußelement (44, 46, 48) dem proximalen Ende des Zweigtransplantats (16) zugeordnet wird, wobei das zweite Anschlußelement (44, 46, 48) konfiguriert wird, um den ersten Anschluß in Eingriff zu nehmen, wenn das proximale Ende des Zweigtransplantats in eine Annäherung mit der Zweigtransplantatöffnung des Haupttransplantats gebracht wird, dadurch gekennzeichnet, daß die eine oder die mehreren Haupttransplantat-Verankerungskomponenten (18) das Haupttransplantat (12) an Stellen auf jeder Seite der anatomischen Zweigröhre halten, wobei die Haupttransplantat-Verankerungskomponenten (18) dadurch wirksam sind, um das Haupttransplantat (12) in einer wesentlich feststehenden Position innerhalb des Hohlraums der anatomischen Röhre zu halten, so daß seine Zweigtransplantatöffnung (14) wesentlich mit der anatomischen Zweigröhre ausgerichtet wird, und das Zweigtransplantat (16) mit dem distalen Ende zuerst durch die Zweigtransplantatöffnung (14) und in die anatomische Zweigröhre vorgeschoben werden kann, wobei das zweite Anschlußelement (44, 46, 48) konfiguriert wird, um selbsttätig reibschlüssig mit dem ersten Anschluß ineinanderzugreifen, um eine Verbindung zwischen dem proximalen Ende des Zweigtransplantats und dem Haupttrans-

plantat zu bilden.

2. System nach Anspruch 1, bei dem wenigstens eine der Komponenten Haupttransplantat (12) und Zweigtransplantat (16) wenigstens teilweise aus gewebtem Polyester hergestellt wird.

3. System nach Anspruch 1, bei dem wenigstens eine der Komponenten Haupttransplantat (12) und Zweigtransplantat (16) wenigstens teilweise aus aufgeschäumtem Polytetrafluorethylen hergestellt wird.

4. System nach Anspruch 1, 2 oder 3, bei dem die Zweigtransplantatöffnung (14) in der Seite des Haupttransplantats (12) geformt wird.

5. System nach Anspruch 1, 2 oder 3, bei dem an einem Ende des Haupttransplantats (12) eine Gabelung geformt wird und die Zweigtransplantatöffnung (14) in einer Seite dieser Gabelung geformt wird.

6. System nach Anspruch 5, bei dem das Haupttransplantat (12) außerdem eine biegsame röhrenförmige Transplantatverlängerung (62) umfaßt, die sich von derjenigen Seite der Gabelung erstreckt, die der Seite gegenüberliegt, in der die Zweigtransplantatöffnung (14) geformt wird.

7. System nach einem der vorhergehenden Ansprüche, bei dem das erste Anschlußelement (40, 42) eine wenigstens teilweise um die Zweigtransplantatöffnung (14) im Haupttransplantat (12) geformte ringförmige Nut umfaßt.

8. System nach einem der vorhergehenden Ansprüche, bei dem die eine oder die mehreren Haupttransplantat-Verankerungskomponenten aus der Gruppe von Transplantat-Verankerungsvorrichtungen und -substanzen ausgewählt werden, die aus folgendem besteht:

wenigstens einem in Radialrichtung ausdehnbaren Stent, der vom Haupttransplantat (12) getrennt ist,

wenigstens einem in Radialrichtung ausdehnbaren Stent, der in das Haupttransplantat (12) integriert ist,

wenigstens einem in Radialrichtung ausdehnbaren Rahmen, der vom Haupttransplantat getrennt ist,

wenigstens einem in Radialrichtung ausdehnbaren Rahmen, der in das Haupttransplantat eingebaut ist,

- wenigstens einem Vorsprung, der wenigstens teilweise durch die Wand der anatomischen Röhre vorsteht,
- einem Klebstoff,
- wenigstens einer Naht
- und wenigstens einer Klammer.
9. System nach einem der vorhergehenden Ansprüche, bei dem der Zweigtransplantat-Abschnitt (16) des Systems außerdem eine Zweigtransplantat-Verankerungskomponente (20) umfaßt, um wenigstens das distale Ende des Zweigtransplantats (16) in Kontakt mit der Wand der anatomischen Zweigröhre zu halten.
10. System nach Anspruch 9, bei dem die Zweigtransplantat-Verankerungskomponente aus der Gruppe von Transplantat-Verankerungsvorrichtungen und -substanzen ausgewählt wird, die aus folgendem besteht:
- wenigstens einem in Radialrichtung ausdehnbaren Stent, der von einer ersten biegsamen Röhre des Zweigtransplantats (16) getrennt ist,
- wenigstens einem in Radialrichtung ausdehnbaren Stent, der in eine erste biegsame Röhre des Zweigtransplantats (16) integriert ist,
- wenigstens einem in Radialrichtung ausdehnbaren Rahmen, der von einer ersten biegsamen Röhre des Zweigtransplantats (16) getrennt ist,
- wenigstens einem in Radialrichtung ausdehnbaren Rahmen, der in eine erste biegsame Röhre des Zweigtransplantats (16) eingebaut ist,
- wenigstens einem Vorsprung, der wenigstens teilweise durch die Wand der anatomischen Röhre vorsteht,
- einem Klebstoff,
- wenigstens einer Naht
- und wenigstens einer Klammer.
11. System nach einem der vorhergehenden Ansprüche, bei dem das zweite Anschlußelement (44, 46, 48) am Zweigtransplantat (16) konfiguriert wird, um reibschlüssig mit dem ersten Anschlußelement (40, 42) des Haupttransplantats (12) ineinanderzugreifen, wenn das Zweigtransplantat (16) aus der Zweigtransplantatöffnung (14) des Haupttransplantats (12) vorgeschoben wird.
12. System nach Anspruch 11, bei dem das erste Anschlußelement (40, 42) am Haupttransplantat (12) eine ringförmige Nut ist und das zweite Anschlußelement (44, 46, 48) am Zweigtransplantat (16) ein ringförmiger Steg ist, der bemessen und konfiguriert wird, um innerhalb der ringförmigen Nut zu sitzen und reibschlüssig mit derselben ineinanderzugreifen.
13. System nach einem der vorhergehenden Ansprüche, bei dem wenigstens ein Abschnitt des Zweigtransplantats (16) verjüngt wird, so daß dessen distales Ende im Durchmesser kleiner ist als dessen proximales Ende.
14. System nach Anspruch 13, bei dem ein Abschnitt des Zweigtransplantats (16), angrenzend an dessen proximales Ende, eine kegelstumpfförmige Konfiguration hat.
15. System nach Anspruch 14, bei dem das erste Anschlußelement (44, 46, 48) am proximalen Ende des kegelstumpfförmigen Segments (44) des Zweigtransplantats (16) geformt wird.
- Revendications**
1. Système de greffe endoluminale d'un conduit anatomique (A) comportant au moins un conduit anatomique dérivé (1A, RA) émanant de celui-ci; ledit système comprenant:
- (a) une greffe tubulaire primaire (12) destinée à être déployée dans le conduit anatomique;
- (i) ladite greffe primaire (12) comportant des première et deuxième extrémités et une lumière la traversant longitudinalement;
- (ii) une ouverture de dérivation de la greffe (14) formée au niveau d'un premier emplacement dans la greffe primaire (12);
- (iii) un premier élément de raccordement (40, 42) formé près de l'ouverture de dérivation de la greffe (14); et
- (iv) un ou plusieurs composants d'ancrage de la greffe primaire (18) destinés à retenir au moins les première et deuxième extrémités de la greffe primaire (12) de sorte qu'elle bute pratiquement contre le conduit anatomique; et

- (b) une greffe de dérivation tubulaire (16) destinée à être déployée dans le conduit anatomique dérivé;
- (i) ladite greffe de dérivation (16) comportant une extrémité proximale, une extrémité distale et une lumière la traversant longitudinalement;
- (ii) un deuxième élément de raccordement (44, 46, 48) étant associé à l'extrémité proximale de la greffe de dérivation (16), ledit deuxième élément de raccordement (44, 46, 48) étant configuré de sorte à s'engager dans ledit premier élément de raccordement, lors du rapprochement de l'extrémité proximale de la greffe de dérivation de l'ouverture de dérivation de la greffe de la greffe primaire, **caractérisé en ce que** un ou plusieurs composants d'ancrage de la greffe primaire (18) retiennent la greffe primaire (12) au niveau d'emplacements situés de chaque côté dudit conduit anatomique de dérivation, lesdits composants d'ancrage de la greffe primaire (18) servant ainsi à retenir la greffe primaire (12) dans une position pratiquement fixe dans la lumière du conduit anatomique, de sorte que son ouverture de dérivation de la greffe (14) est pratiquement alignée avec ledit conduit anatomique de dérivation; ladite greffe de dérivation (16) pouvant être avancée à partir de l'extrémité distale à travers l'ouverture de dérivation de la greffe (14) et dans le conduit anatomique de dérivation; ledit deuxième élément de raccordement (44, 46, 48) étant configuré de sorte à s'engager automatiquement par frottement dans ledit premier élément de raccordement pour établir une connexion entre l'extrémité proximale de la greffe de dérivation et la greffe primaire.
2. Système selon la revendication 1, dans lequel au moins ladite greffe primaire (12) ou ladite greffe de dérivation (16) est composée au moins partiellement de polyester tissé.
 3. Système selon la revendication 1, dans lequel au moins ladite greffe primaire (12) ou ladite greffe de dérivation (16) est composée au moins partiellement de polytétrafluoréthylène expansé.
 4. Système selon les revendications 1, 2 ou 3, dans lequel l'ouverture de dérivation de la greffe (14) est formée dans le côté de la greffe primaire (12).
 5. Système selon les revendications 1, 2 ou 3, dans lequel une bifurcation est formée au niveau d'une extrémité de la greffe primaire (12), ladite ouverture de dérivation de la greffe (14) étant formée dans un côté de cette bifurcation.
 6. Système selon la revendication 5, dans lequel la greffe primaire (12) comprend en outre une extension de greffe tubulaire pliable (62) s'étendant à partir du côté de la bifurcation opposée au côté dans lequel est formée l'ouverture de dérivation de la greffe (14).
 7. Système selon l'une quelconque des revendications précédentes, dans lequel le premier élément de raccordement (40, 42) comprend une rainure annulaire formée au moins partiellement autour de l'ouverture de dérivation de la greffe (14) dans la greffe primaire (12).
 8. Système selon l'une quelconque des revendications précédentes, dans lequel l'un ou les plusieurs composants d'ancrage de la greffe primaire sont sélectionnés dans le groupe comprenant un dispositif d'ancrage de la greffe et des substances constituées de:
 - au moins un stent à dilatation radiale séparé de la greffe primaire (12);
 - au moins un stent à dilatation radiale intégré dans la greffe primaire (12);
 - au moins un cadre à dilatation radiale séparé de la greffe primaire;
 - au moins un cadre à dilatation radiale incorporé dans la greffe primaire;
 - au moins une saillie débordant au moins partiellement à travers la paroi du conduit anatomique;
 - et d'un adhésif;
 - et d'au moins une suture;
 - et
 - d'au moins une agrafe.
 9. Système selon l'une quelconque des revendications précédentes, dans lequel la partie de la greffe de dérivation (16) dudit système comprend un composant d'ancrage de la greffe de dérivation (20) destiné à retenir au moins l'extrémité distale de la greffe de dérivation (16) en contact avec la paroi du conduit anatomique de dérivation.
 10. Système selon la revendication 9, dans lequel le composant d'ancrage de la greffe de dérivation est sélectionné du groupe comprenant un dispositif d'ancrage de la greffe et des substances constituées de:
 - au moins un stent à dilatation radiale séparé d'un premier tube pliable de la greffe de dériva-

- tion (16);
- au moins un stent à dilatation radiale intégré dans un premier tube pliable de la greffe de dérivation (16); 5
- au moins un cadre à dilatation radiale séparé d'un premier tube pliable de la greffe de dérivation (16); 10
- au moins un cadre à dilatation radiale incorporé dans un premier tube pliable de la greffe de dérivation (16);
- au moins une saillie débordant au moins partiellement à travers la paroi du conduit anatomique; 15
- un adhésif; 20
- au moins une suture;
- et au moins une agrafe.
11. Système selon l'une quelconque des revendications précédentes, dans lequel le deuxième élément de raccordement (44, 46, 48) sur la greffe de dérivation (16) est configuré de sorte à s'engager par frottement dans le premier élément de raccordement (40, 42) de la greffe primaire (12) lors de la sortie de la greffe de dérivation (16) de l'ouverture de dérivation de la greffe (14) de la greffe primaire (12). 25 30
12. Système selon la revendication 11, dans lequel le premier élément de raccordement (40, 42) sur la greffe primaire (12) est constitué par une rainure annulaire, le deuxième élément de raccordement (44, 46, 48) sur la greffe de dérivation (16) étant constitué par une nervure annulaire dimensionnée et configurée de sorte à être logée dans ladite rainure annulaire et à s'engager par frottement dans celle-ci. 35 40
13. Système selon l'une quelconque des revendications précédentes, dans lequel au moins une partie de la greffe de dérivation (16) est effilée, de sorte que son extrémité distale a un diamètre inférieur à celui de l'extrémité proximale. 45 50
14. Système selon la revendication 13, dans lequel une partie de la greffe de dérivation (16) a une configuration en tronc de cône près de son extrémité proximale. 55
15. Système selon la revendication 14, dans lequel le premier élément de raccordement (44, 46, 48) est formé sur l'extrémité proximale du segment en tronc

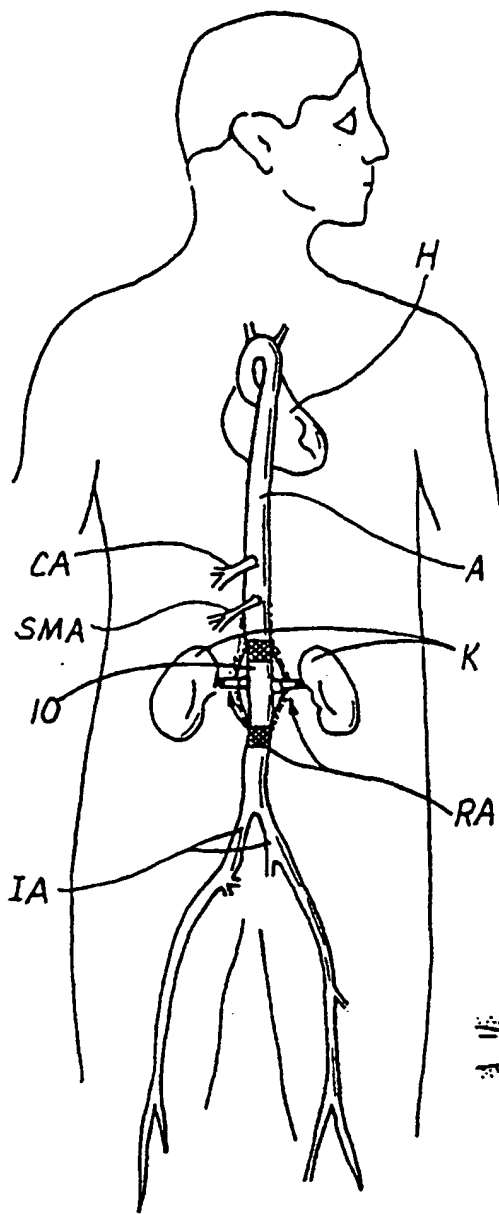


Fig. 1

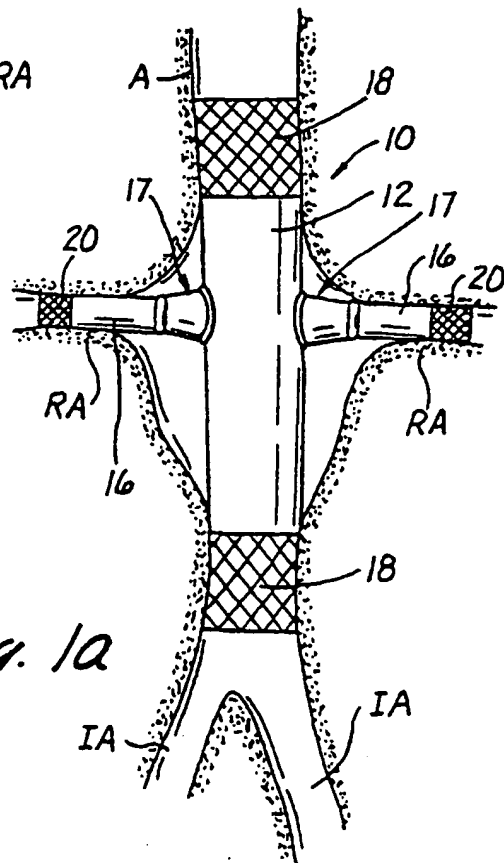
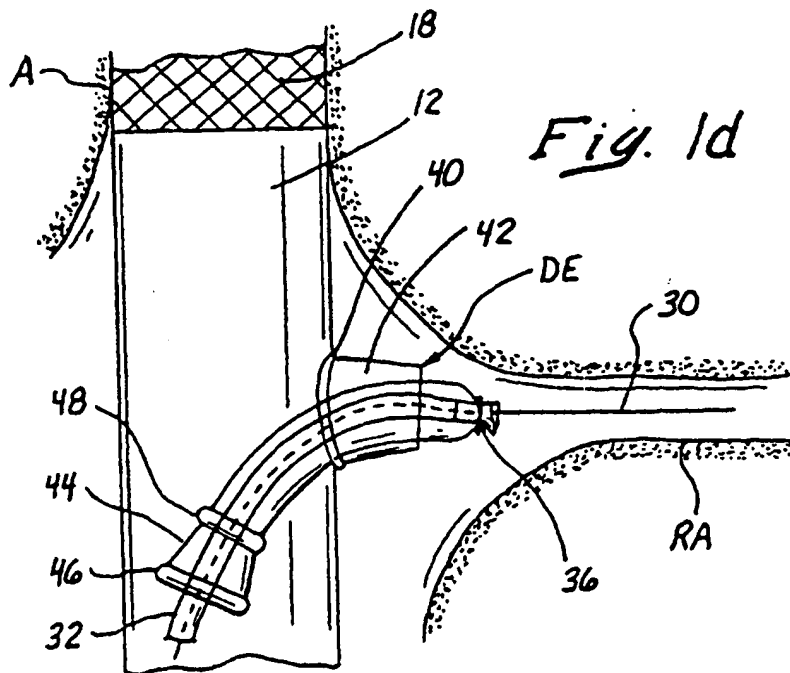
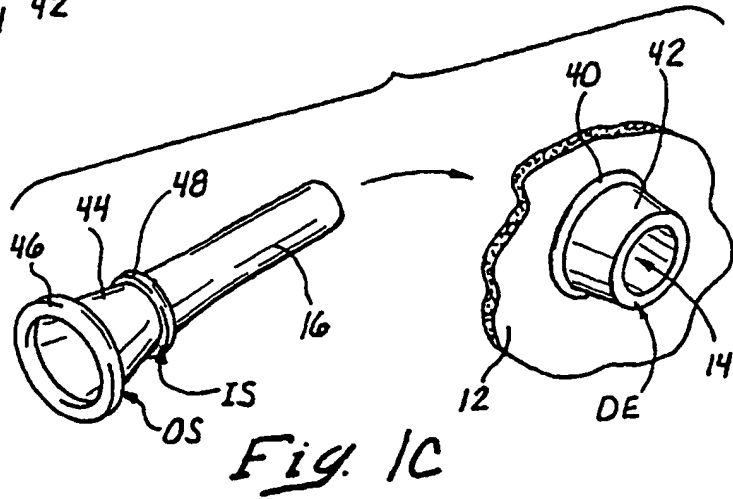
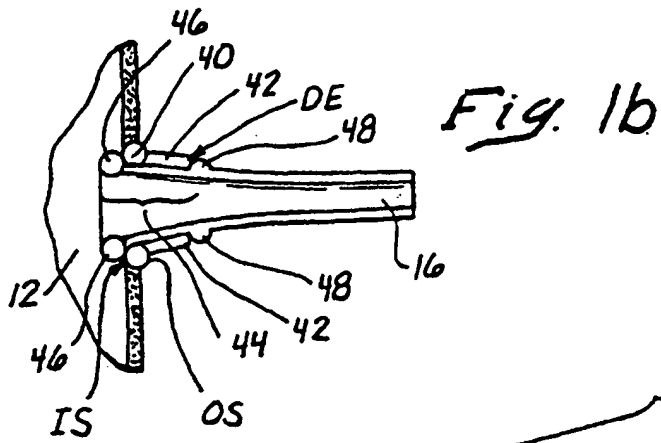
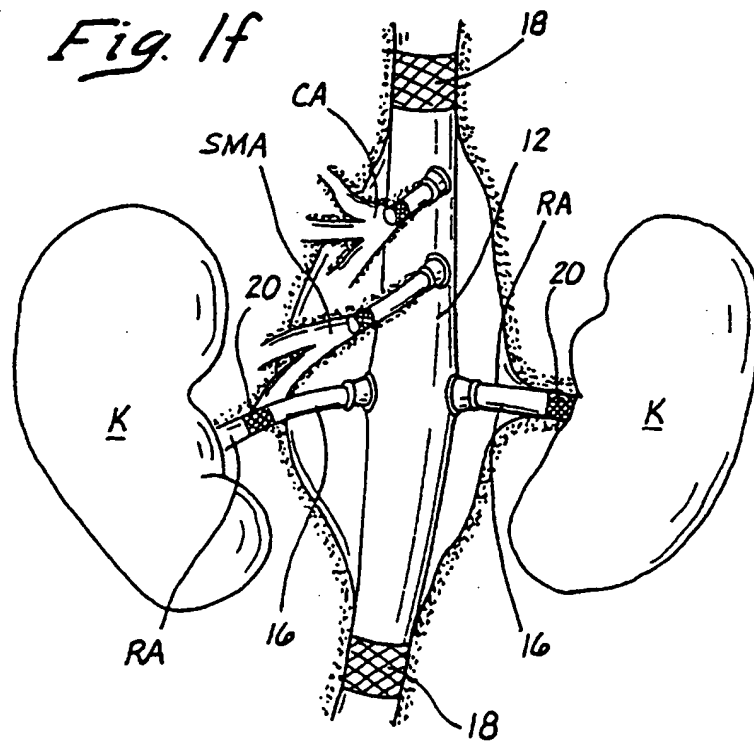
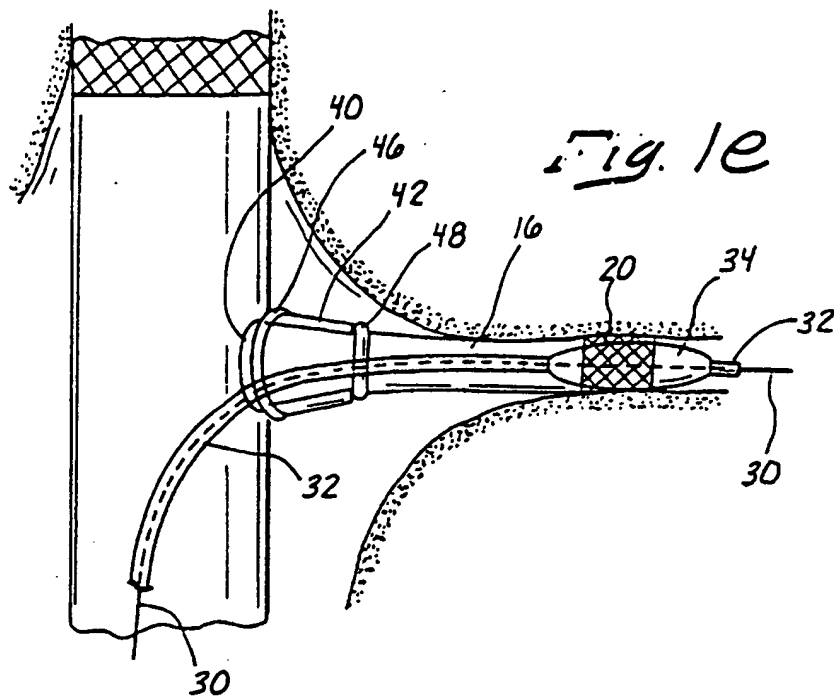
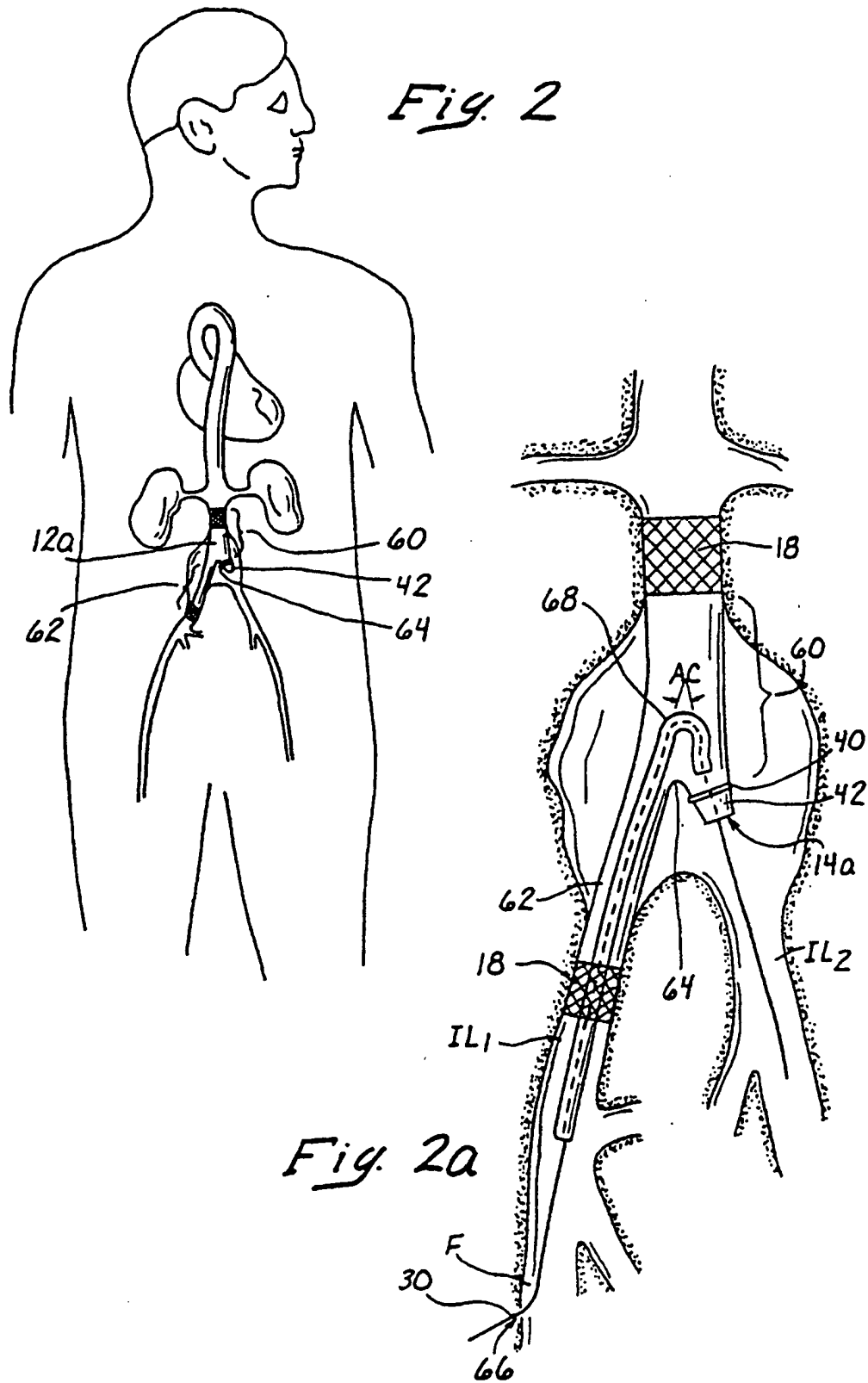
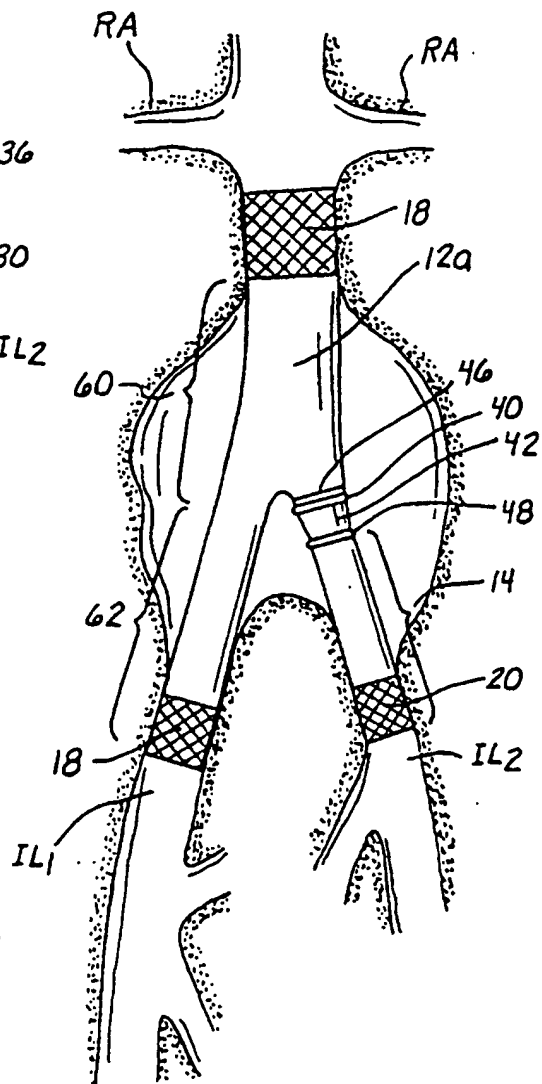
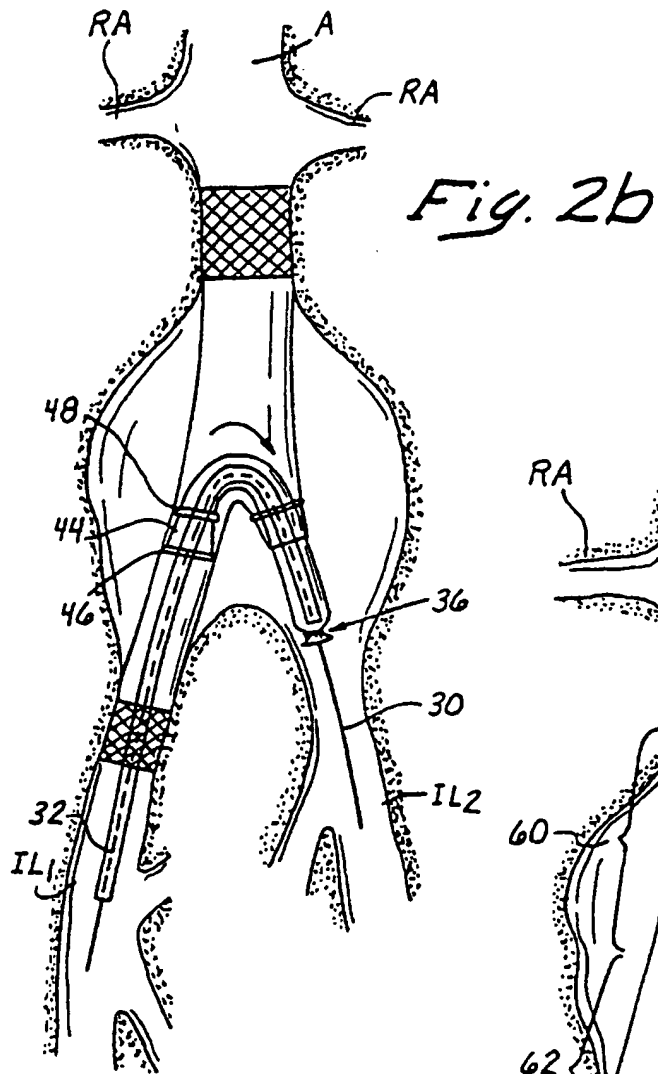


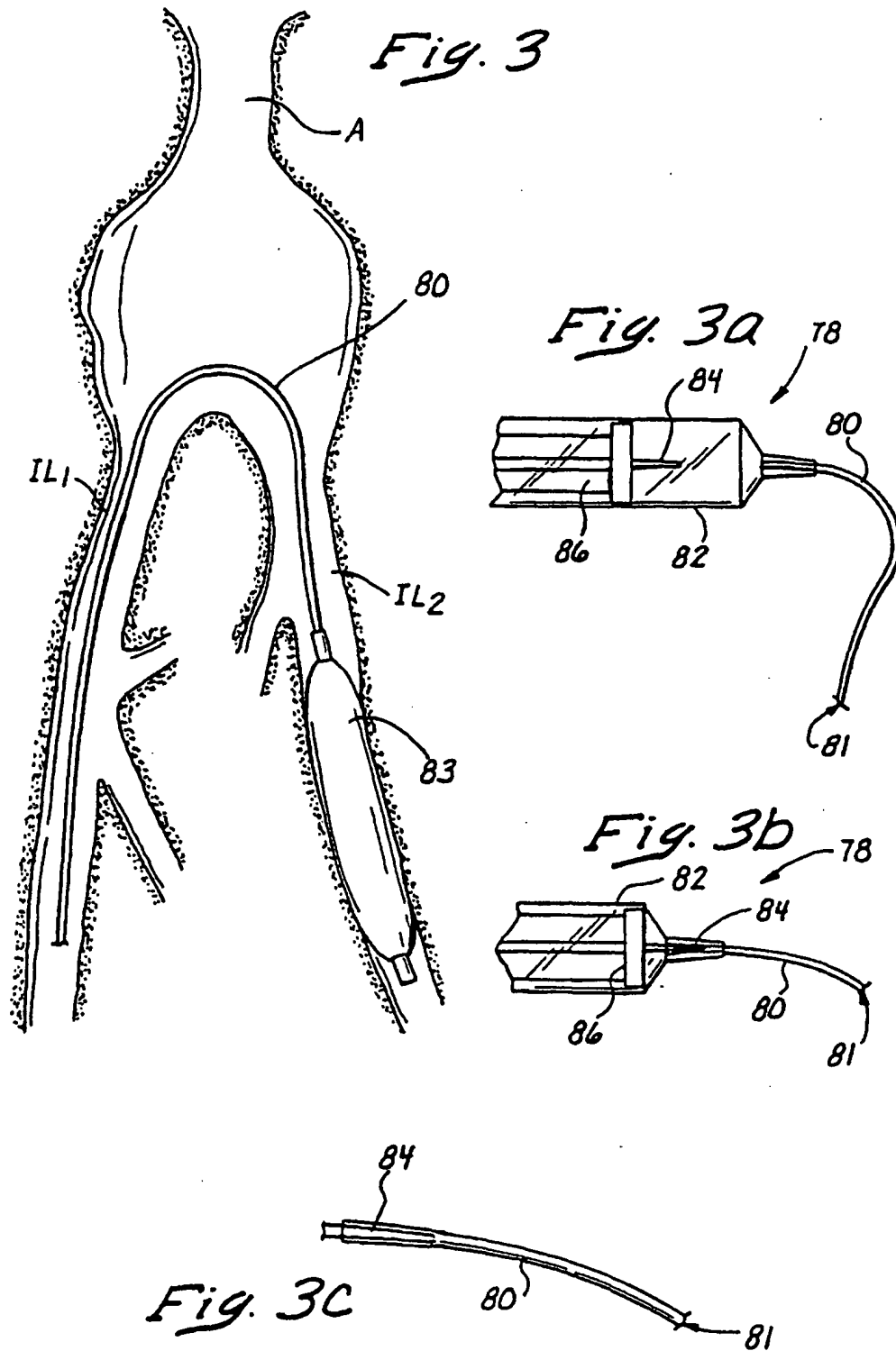
Fig. 1a











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